Refine Search

Search Results -

Term	Documents
(5 NOT 3).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	12
(L5 NOT L3).PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD.	12

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Database:

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Search History

DATE: Friday, April 15, 2005 Printable Copy Create Case

Set Name side by side	Query	Hit Count	<u>Set</u> <u>Name</u> result set
<i>DB=PGPB</i> <i>OP=AND</i>	B,USPT,USOC,EPAB,JPAB,DWPI,TDBD; THES=ASSIGNEE; PLU	VR = YES;	
<u>L6</u>	L5 not L3	12	<u>L6</u>
<u>L5</u>	((modified or mutant) same (lambda adj integrase))	18	<u>L5</u>
<u>L4</u>	L2 not L3	38	<u>L4</u>
<u>L3</u>	L2 and (attB or attP or attL or attR)	7	<u>L3</u>
<u>L2</u>	((modified or mutant) adj Int) or (Int-h or Int-h/218)	45	<u>L2</u>
<u>L1</u>	Droge-Peter.in.	2	<u>L1</u>

END OF SEARCH HISTORY

Set



PALM INTRANET

Day: Friday Date: 4/15/2005

Time: 11:16:26

Inventor Name Search

Enter the first few letters of the Inventor's Last Name. Additionally, enter the first few letters of the Inventor's First name.

Last Name	First Name	
Droge	Peter	Search

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***CorpTech (559)
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KWIC is set to 50.
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 * * *
File 1:ERIC 1966-2004/Jul 21
      (c) format only 2004 The Dialog Corporation
 *File
         1: Updates suspended by ERIC until
Q2, 2005
      Set Items Description
      --- ---- ------
Cost is in DialUnits
B 155, 159, 5, 73
       15apr05 12:24:05 User259876 Session D741.1
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     $0.80 Estimated cost File1
     $0.06 INTERNET
     $0.86 Estimated cost this search
     $0.86 Estimated total session cost 0.228 DialUnits
SYSTEM:OS - DIALOG OneSearch
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         (c) format only 2005 The Dialog Corp.
  File 159: Cancerlit 1975-2002/Oct
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         5:Biosis Previews(R) 1969-2005/Apr W2
         (c) 2005 BIOSIS
  File 73:EMBASE 1974-2005/Apr W2
         (c) 2005 Elsevier Science B.V.
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S (MODIFIED OR MUTANT) (S) (LAMBDA (W) INTEGRASE)
          532363 MODIFIED
          527565 MUTANT
           83776 LAMBDA
            6950 INTEGRASE
      S1
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?
S (INT-H) OR (INT-H/218)
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               1 INT-H/218
              3 (INT-H) OR (INT-H/218)
      S2
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RD S2
...completed examining records
              2 RD S2 (unique items)
?
T S3/3, K/ALL
  3/3, K/1
             (Item 1 from file: 155)
DIALOG(R) File 155:MEDLINE(R)
(c) format only 2005 The Dialog Corp. All rts. reserv.
10882829
           PMID: 7874687
 The overexpression of int-5/Aromatase, a novel MMTV integration locus
 gene, is responsible for D2 mammary tumor cell proliferation.
  Tekmal R R; Durgam V R
  Department of Obstetrics and Gynecology,
                                                University of Texas Health
Science Center at San Antonio 78284-7836.
          letters (IRELAND)
  Cancer
                              Jan
                                     27
                                          1995,
                                                 88
                                                       (2)
                                                             p147-55,
0304-3835
          Journal Code: 7600053
  Contract/Grant No.: P30 CA 54174; CA; NCI; R29 CA57559; CA; NCI
  Publishing Model Print
  Document type: Journal Article
  Languages: ENGLISH
 Main Citation Owner: NLM
  Record type: MEDLINE; Completed
  Gene Symbol: MMTV; P450; int-5; int-H
  3/3, K/2
              (Item 1 from file: 5)
DIALOG(R) File
                5:Biosis Previews(R)
(c) 2005 BIOSIS. All rts. reserv.
0012039800
            BIOSIS NO.: 199900299460
```

```
Alterations in the directionality of lambda site-specific recombination
 catalyzed by mutant integrases in vivo
AUTHOR: Christ Nicole; Droege Peter (Reprint)
AUTHOR ADDRESS: Institute of Genetics, University of Cologne, Weyertal 121,
  D-50931, Cologne, Germany**Germany
JOURNAL: Journal of Molecular Biology 288 (5): p825-836 May 21, 1999 1999
MEDIUM: print
ISSN: 0022-2836
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
DESCRIPTORS:
  CHEMICALS & BIOCHEMICALS:
                              ... Int-h/218
... Int-h
?
Set
        Items
                Description
S1
           69
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S2
            3
                (INT-H) OR (INT-H/218)
S3
            2
                RD S2 (unique items)
S S1 NOT PY>2000
              69 S1
         6706539 PY>2000
              41 S1 NOT PY>2000
?
...completed examining records
      S5
             14 RD (unique items)
T S5/3, K/ALL
  5/3, K/1
              (Item 1 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2005 The Dialog Corp. All rts. reserv.
13304409
          PMID: 10075917
 X-ray structure of T4 endonuclease VII: a DNA junction resolvase with a
 novel fold and unusual domain-swapped dimer architecture.
  Raaijmakers H; Vix O; Toro I; Golz S; Kemper B; Suck D
  Structural
             Biology Programme,
                                     EMBL,
                                             Meyerhofstrasse
Heidelberg, Germany.
  EMBO journal (ENGLAND)
                           Mar 15 1999, 18 (6) p1447-58, ISSN 0261-4189
Journal Code: 8208664
  Publishing Model Print
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: MEDLINE; Completed
  ... containing a zinc ion tetrahedrally coordinated to four cysteines,
does not resemble any of the known junction-resolving enzymes, including
the Escherichia coli RuvC and lambda integrase -type recombinases. The
S-shaped dimer has two 'binding bays' separated by approximately 25 A which
```

are lined by positively charged residues and contain near...

5/3,K/2 (Item 2 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

13274883 PMID: 9931245

DNA-sequence asymmetry directs the alignment of recombination sites in the FLP synaptic complex.

Huffman K E; Levene S D

Department of Molecular and Cell Biology, The University of Texas at Dallas, Richardson, TX, PO Box 830688, USA.

Journal of molecular biology (ENGLAND) Feb 12 1999, 286 (1) p1-13, ISSN 0022-2836 Journal Code: 2985088R

Contract/Grant No.: GM47898; GM; NIGMS

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... FLP synaptic complexes examined contain the two FRTs aligned in parallel. This strong preference for parallel site alignment stands in contrast with prevailing models for lambda integrase -class recombination systems, which postulate antiparallel site alignment, and results from biophysical studies on synthetic, immobile four-way DNA junctions. Our results show that the...

5/3,K/3 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

12980970 PMID: 10939248

Genetic characterization of gram-positive homologs of the XerCD site-specific recombinases.

Chalker A F; Lupas A; Ingraham K; So C Y; Lunsford R D; Li T; Bryant A; Holmes D J; Marra A; Pearson S C; Ray J; Burnham M K; Palmer L M; Biswas S; Zalacain M

SmithKline Beecham Pharmaceuticals, Collegeville, PA 19426-0989, USA. Alison F Chalker@sbphrd.com

Journal of molecular microbiology and biotechnology (ENGLAND) Apr 2000, 2 (2) p225-33, ISSN 1464-1801 Journal Code: 100892561

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... the S. aureus xerD gene appears to be absolutely required for viability, and may therefore be the first example of an essential gene of the lambda integrase family. In contrast, phylogenetic and conservation pattern analysis show that the S. pneumoniae gene products are more closely related to phage integrases than to XerCD...

5/3,K/4 (Item 4 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

12800974 PMID: 10698624

Site-specific recombination in human cells catalyzed by phage lambda integrase mutants.

Lorbach E; Christ N; Schwikardi M; Droge P

Institute of Genetics, University of Cologne, Weyertal 121, Cologne, D-50931, Germany.

Journal of molecular biology (ENGLAND) Mar 10 2000, 296 (5) p1175-81 ISSN 0022-2836 Journal Code: 2985088R

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

...to Cre and FLP, however, wild-type Int requires accessory proteins and DNA supercoiling of target sites to catalyze recombination. Here, we show that two mutant Int proteins, Int-h (E174 K) and its derivative Int-h/218 (E174 K/E218 K), which do not require accessory factors, are proficient to . . .

5/3,K/5 (Item 5 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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12784278 PMID: 10713132

Roles of topoisomerases in maintaining steady-state DNA supercoiling in Escherichia coli.

Zechiedrich E L; Khodursky A B; Bachellier S; Schneider R; Chen D; Lilley D M; Cozzarelli N R

Department of Molecular Virology and Microbiology, Baylor College of Medicine, Houston, Texas 77030, USA.

Journal of biological chemistry (UNITED STATES) Mar 17 2000, p8103-13, ISSN 0021-9258 Journal Code: 2985121R

Contract/Grant No.: ESO1890; ES; NIEHS; GM 31657; GM; NIGMS

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... become hyper-negatively supercoiled (sigma = -0.09), greatly stimulating transcription from the supercoiling sensitive leu-500 promoter and increasing the number of supercoils trapped by lambda site-specific recombination.

5/3,K/6 (Item 6 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

12455195 PMID: 9765577

Point mutations in the integron integrase IntI1 that affect recombination and/or substrate recognition.

Gravel A; Messier N; Roy P H

Centre de Recherche en Infectiologie, Centre Hospitalier de l'Universite Laval and Departement de Biochimie, Faculte des Sciences et de Genie, Universite Laval, Sainte-Foy, Quebec, Canada.

Journal of bacteriology (UNITED STATES) Oct 1998, 180 (20) p5437-42, Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... lower than that obtained with the wild-type MBP-IntI1. We have also made two proteins with mutations of the tyrosine residue (Y312), and both mutant proteins are similar to the wild-type fusion protein in their DNA-binding capacity but are unable to catalyze in vivo recombination.

5/3,K/7 (Item 7 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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12364806 PMID: 9675854

Cloning and characterisation of the Proteus mirabilis xerD gene.

Villion M; Szatmari G

Departement de Microbiologie et Immunologie, Universite de Montreal, Que., Canada.

FEMS microbiology letters (NETHERLANDS) Jul 1 1998, 164 (1) p83-90, ISSN 0378-1097 Journal Code: 7705721

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... methods. This gene encodes a tyrosine recombinase which is highly similar to the E. coli XerD recombinase, is capable of complementing an E. coli xerD mutant , and displays sequence-specific DNA binding activity.

5/3, K/8 (Item 8 from file: 155)

DIALOG(R)File 155:MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

12310999 PMID: 9618537

A site-specific recombinase is required for competitive root colonization by Pseudomonas fluorescens WCS365.

Dekkers L C; Phoelich C C; van der Fits L; Lugtenberg B J

Leiden University, Institute of Molecular Plant Sciences, Clusius Laboratory, Wassenaarseweg 64, 2333AL Leiden, The Netherlands.

Proceedings of the National Academy of Sciences of the United States of America (UNITED STATES) Jun 9 1998, 95 (12) p7051-6, ISSN 0027-8424 Journal Code: 7505876

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... sss gene homologue rather than orf240 is crucial for colonization. xerC in Escherichia coli and sss in Pseudomonas aeruginosa encode proteins that belong to the lambda integrase family of site-specific recombinases, which play a role in phase variation caused by DNA rearrangements. The function of the xerC/sss homologue in colonization is discussed in terms of genetic rearrangements involved in the generation of

different phenotypes, thereby allowing a bacterial population to occupy various habitats. **Mutant** PCL1233 is assumed to be locked in a phenotype that is not well suited to compete for colonization in the rhizosphere. Thus we show the...

5/3,K/9 (Item 9 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

12268821 PMID: 9571022

Recognition of core binding sites by bacteriophage integrases.

Dorgai L; Sloan S; Weisberg R A

Laboratory of Molecular Genetics, National Institute of Child Health and Human Development, Bethesda, MD 20892, USA.

Journal of molecular biology (ENGLAND) Apr 17 1998, 277 (5) p1059-70

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

...on recombination, suggesting that integrase does not recognize each of the extended core binding sites in the same way. Finally, substitution at several positions in **lambda** integrase with the corresponding HK022-specific amino acids prevents recombination of lambda attachment sites, and this defect can be suppressed in an allele-specific manner by...

5/3,K/10 (Item 10 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

(c) format only 2005 The Dialog Corp. All rts. reserv.

10964424 PMID: 7744017

Xer site-specific recombination in vitro.

Arciszewska L K; Sherratt D J

Department of Biochemistry, University of Oxford, UK.

EMBO journal (ENGLAND) May 1 1995, 14 (9) p2112-20, ISSN 0261-4189

Journal Code: 8208664
Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... corresponding to the conversion of the Holliday junction intermediate back to the substrate, has been observed. Recombination reactions using XerC and XerD derivatives that are **mutant** in their presumptive catalytic residues, or are maltose-binding fusion recombinase derivatives, have demonstrated that this pair of strand exchanges is catalysed by XerC. The

5/3,K/11 (Item 11 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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10933558 PMID: 7715441

The sss gene product, which affects pyoverdin production in Pseudomonas

aeruginosa 7NSK2, is a site-specific recombinase.

Hofte M; Dong Q; Kourambas S; Krishnapillai V; Sherratt D; Mergeay M Laboratory of Phytopathology, University of Gent, Belgium.

Molecular microbiology (ENGLAND) Dec 1994, 14 (5) p1011-20, ISSN

0950-382X Journal Code: 8712028

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... this mutant was cloned and sequenced. Its protein sequence showed 50% identity to the XerC protein of Escherichia coli, which is a member of the lambda integrase family of site-specific recombinases. An open reading frame was found upstream of sss whose protein sequence showed strong identity to DapF, the diaminopimelate epimerase...

... a multicistronic unit that also contains dapF. The sss gene of P. aeruginosa could restore site-specific recombination at cer in an E. coli xerC mutant and the E. coli xerC gene could complement a genomic sss mutation in P. aeruginosa.

5/3,K/12 (Item 12 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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08905232 PMID: 2155203

Genetic analysis of bacteriophage lambda integrase interactions with arm-type attachment site sequences.

Lee E C; Gumport R I; Gardner J F

Department of Microbiology, University of Illinois, Urbana 61801.

Journal of bacteriology (UNITED STATES) Mar 1990, 172 (3) p1529-38,

Contract/Grant No.: GM28717; GM; NIGMS

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... by binding to the same arm-type sites. Mutations in the P'123 or P'1 sites that impair Int binding were isolated by selecting mutant phages that express antirepressor in the presence of Int. DNA sequence analyses showed that most of the mutants in the challenge phages carrying the P...

5/3,K/13 (Item 13 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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08265612 PMID: 2836392

Suicide recombination substrates yield covalent lambda integrase-DNA complexes and lead to identification of the active site tyrosine.

Pargellis C A; Nunes-Duby S E; de Vargas L M; Landy A

Division of Biology and Medicine, Brown University, Providence, Rhode Island 02912.

Journal of biological chemistry (UNITED STATES) Jun 5 1988, 263 (16) p7678-85, ISSN 0021-9258 Journal Code: 2985121R

Contract/Grant No.: GM33928; GM; NIGMS

Publishing Model Print

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Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: MEDLINE; Completed
  ... lambda integrase protein. The tyrosine residue at position 342 is
shown to form a covalent bond with DNA at the sites of strand exchange. A
  mutant integrase in which this tyrosine is changed to phenylalanine is
devoid of both topoisomerase and recombinase activity but still binds to
both core- and arm...
  5/3,K/14
               (Item 1 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
(c) 2005 BIOSIS. All rts. reserv.
             BIOSIS NO.: 199800510299
0011716052
 Point mutations in the integron intergrasse IntI1 that affect recombination
 and/or substrate recognition
AUTHOR: Gravel Annie; Messier Nancy; Roy Paul H (Reprint)
AUTHOR ADDRESS: Cent. Recherche Infectiol., CHUL, Local RC-709, 2705 Boul.
  Laurier, Sainte-Foy, PQ G1V 4G2, Canada**Canada
JOURNAL: Journal of Bacteriology 180 (20): p5437-5442 Oct., 1998 1998
MEDIUM: print
ISSN: 0021-9193
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
... ABSTRACT: lower than that obtained with the wild-type MmBP-IntI1. We
  have also made two proteins with mutations of the tyrosine residue
  (Y312), and both mutant proteins are similar to the wild-type fusion
  protein in their DNA-binding capacity but are unable to catalyze in vivo
  recombination.
?
        Items
               Description
S1
           69
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S2
           3
              (INT-H) OR (INT-H/218)
S3
           2 RD S2 (unique items)
S4
           41
               S1 NOT PY>2000
S5
          14
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S (LAMBDA (W) INTEGRASE) (S) (HUMAN OR MAMMALIAN OR EUKARYOTIC OR MOUSE)
          83776 LAMBDA
            6950 INTEGRASE
        14542331 HUMAN
          354798 MAMMALIAN
          94567 EUKARYOTIC
         1699644 MOUSE
             25 (LAMBDA (W) INTEGRASE) (S) (HUMAN OR MAMMALIAN OR
                EUKARYOTIC OR MOUSE)
?
S S6 AND (ATTB OR ATTP OR ATTL OR ATTR)
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             633 ATTB
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878 ATTP
             292 ATTL
             485 ATTR
      s7
              11 S6 AND (ATTB OR ATTP OR ATTL OR ATTR)
RD
...completed examining records
               3 RD (unique items)
T S8/3, K/ALL
  8/3,K/1
              (Item 1 from file: 155)
DIALOG(R) File 155: MEDLINE(R)
(c) format only 2005 The Dialog Corp. All rts. reserv.
           PMID: 14687564
 Phage integrases: biology and applications.
  Groth Amy C; Calos Michele P
  Department of Genetics, Stanford University School of Medicine, Stanford,
CA 94305-5120, USA.
  Journal of molecular biology (England)
                                           Jan 16 2004, 335
                                                              (3) p667-78,
ISSN 0022-2836
               Journal Code: 2985088R
  Contract/Grant No.: DK58187; DK; NIDDK; HL68112; HL; NHLBI
  Publishing Model Print
  Document type: Journal Article; Review; Review, Tutorial
  Languages: ENGLISH
 Main Citation Owner: NLM
  Record type: MEDLINE; Completed
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Phage integrases are enzymes that mediate unidirectional site-specific recombination between two DNA recognition sequences, the phage attachment site, attP , and the bacterial attachment site, attB . Integrases may be grouped into two major families, the tyrosine recombinases and the serine recombinases, based on their mode of catalysis. Tyrosine family integrases, such as lambda integrase, utilize a catalytic tyrosine to mediate strand cleavage, tend to recognize longer attP sequences, and require other proteins encoded by the phage or the host bacteria. Phage integrases from the serine family are larger, use a catalytic serine for strand cleavage, recognize shorter attP sequences, and do not require host cofactors. Phage integrases mediate efficient site-specific recombination between two different sequences that are relatively short, yet long enough to be specific on a genomic scale. These properties give phage integrases growing importance for the genetic manipulation of living eukaryotic cells, especially those with large genomes such as mammals and most plants, for which there are few tools for precise manipulation of the genome. Integrases of the serine family have been shown to work efficiently in mammalian cells, mediating efficient integration at introduced att sites or native sequences that have partial identity to att sites. This reaction has applications in areas such...

```
8/3,K/2 (Item 2 from file: 155)
DIALOG(R)File 155:MEDLINE(R)
(c) format only 2005 The Dialog Corp. All rts. reserv.

12800974 PMID: 10698624
   Site-specific recombination in human cells catalyzed by phage lambda integrase mutants.
```

Lorbach E; Christ N; Schwikardi M; Droge P Institute of Genetics, University of Cologne, Weyertal 121, Cologne,

D-50931, Germany.

Journal of molecular biology (ENGLAND) Mar 10 2000, 296 (5) p1175-81 ISSN 0022-2836 Journal Code: 2985088R

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Site-specific recombination in human cells catalyzed by phage lambda integrase mutants.

... 218 (E174 K/E218 K), which do not require accessory factors, are proficient to perform intramolecular integrative and excisive recombination in co-transfection assays inside human cells. Intramolecular integrative recombination is also detectable by Southern analysis in human reporter cell lines harboring target sites attB and attP as stable genomic sequences. Recombination by wild-type Int, however, is not detectable by this method. The latter result implies that eukaryotic co-factors, which could functionally replace the prokaryotic ones normally required for wild-type Int, are most likely not present in human cells. Copyright 2000 Academic Press.

8/3,K/3 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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12168317 PMID: 9472558

A new DNA vehicle for nonviral gene delivery: supercoiled minicircle.

Darquet A M; Cameron B; Wils P; Scherman D; Crouzet J

UMR 133 CNRS/Rhone-Poulenc Rorer, Vitry sur Seine, France.

Gene therapy (ENGLAND) Dec 1997, 4 (12) p1341-9, ISSN 0969-7128

Journal Code: 9421525
Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

... a high level of biological containment. They are obtained in E. coli by att site-specific recombination mediated by the phage lambda integrase. The desired **eukaryotic** expression cassette, bounded by the lambda **attP** and **attB** sites was cloned on a recombinant plasmid. The expression cassette was excised in vivo after thermoinduction of the integrase gene leading to the formation of...

```
Items Description
Set
S1
           69
                (MODIFIED OR MUTANT) (S) (LAMBDA (W) INTEGRASE)
S2
            3
                (INT-H) OR (INT-H/218)
s3
           2 RD S2 (unique items)
S4
           41 S1 NOT PY>2000
S5
           14
                RD (unique items)
S6
           25
                (LAMBDA (W) INTEGRASE) (S) (HUMAN OR MAMMALIAN OR EUKARYOT-
            IC OR MOUSE)
s7
                S6 AND (ATTB OR ATTP OR ATTL OR ATTR)
           11
S8
                RD (unique items)
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COST
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           $1.76 0.551 DialUnits File155
              $3.57 17 Type(s) in Format 3
           $3.57 17 Types
    $5.33 Estimated cost File155
           $0.48
                  0.163 DialUnits File159
    $0.48 Estimated cost File159
           $4.69 0.816 DialUnits File5
              $4.00 2 Type(s) in Format 3
           $4.00 2 Types
    $8.69 Estimated cost File5
           $7.02 0.660 DialUnits File73
    $7.02 Estimated cost File73
           OneSearch, 4 files, 2.189 DialUnits FileOS
    $2.40 INTERNET
   $23.92 Estimated cost this search
   $24.78 Estimated total session cost 2.418 DialUnits
```

Return to logon page!